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Virulence and antibiotic resistance of isolates of *Klebsiella pneumoniae* in newborns with localized and generalized forms of infection

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Abstract

© The authors team, 2018. All Rights Reserved. Objective. To study the effect of virulence and antibiotic sensitivity of *K. pneumoniae* on the course and outcome of localized and generalized forms of infection in newborns. The authors studied 25 samples of *K. pneumoniae* isolated from the blood (12 isolates) and feces (13 isolates) of the children with various forms of neonatal infection. Group 1 consisted of 12 children with bacteriologically proven neonatal sepsis, *K. pneumoniae* was isolated of their blood. Group 2 included 13 children with localized bacterial infection in the form of pneumonia, *K. pneumoniae* was isolated from their feces. The PCR method was used to determine the virulence factors of the isolates of *K. pneumoniae*-*rmpA*, aerobactin and colibactin. The sensitivity of *K. pneumoniae* to antibiotics was determined by the Kirby-Bauer method. The double disk method was used to determine the ability of *K. pneumoniae* to produce extended-spectrum β -lactamases (ESBL). Results. In Group 1 the isolates of *K. pneumoniae* produced ESBL in 8 children out of 12. The bacteria were sensitive to meropenem, amikacin and ciprofloxacin in 4 cases. One child demonstrated resistance to meropenem. The remaining 4 isolates were sensitive to the third-generation cephalosporins protected by aminopenicillins, amikacin, meropenem and ciprofloxacin. The *rmpA* gene was determined in the *K. pneumoniae* isolates in 6 children. The "string-test" of these colonies of *K. pneumoniae* in all cases gave a positive result. The genes of siderophores, aerobactin and colibactin were found in 3 isolates. Aerobactin and colibactin produced only *rmpA*-bearing strains. 3 isolates (23%) of *K. pneumoniae* produced ESBL in Group 2. In 8 out of 13 cases there was *rmpA*-gene and genes of aerobactin and colibactin in 11 and 7 cases accordingly. The "string-test" was positive in 8 cases, and there were only *rmpA*-positive bacteria. Siderophores were detected both in *rmpA*-positive and *rmpA*-negative isolates. The microbes produced BLBR and were *rmpA*-positive in 2 children. In one case, the isolates had neither the characteristic virulence factors, nor BLBR. Conclusion. The risk of developing localized and generalized forms of neonatal *klebsiella* infection is largely determined by microbiological features of the microorganism, its resistance and virulence. We observed clinical variants of the disease caused by *K. pneumoniae* which simultaneously had two properties: high aggressiveness and resistance to antibiotic therapy.

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Keywords

Antibiotic resistance, Children, *Klebsiella pneumoniae*, Neonatal sepsis, Virulence

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